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## **Claims**

- A method for the treatment of a cancer comprising co-administering
  an anti-tumor antibody and a cytokine to a subject in need thereof,
  wherein the cytokine is administered continuously or repeatedly in a low-dose form.
- 2. A method for the treatment of a cancer comprising coadministering an anti-tumor antibody and a cytokine to a subject in need thereof, wherein the method comprises:
  - (a) a first treatment stage comprising administering a low-dose cytokine, and
  - (b) a second treatment stage comprising co-administering an antitumor antibody and a low-dose cytokine.
  - 3. The method of claim 1 or 2, wherein the low-dose cytokine comprises a dose which is pharmaceutically effective in the substantial absence of NIC CTC toxicity grade 3 or higher.
  - 4. The method according to any one of claims 1-3 comprising a daily administration of a low-dose cytokine.
  - The method of any one of claims 1-4 wherein the cytokine is selected from interleukins and interferons.
    - 6. The method of claim 5 wherein the cytokine is IL-2.
- 7. The method of claim 6 wherein the dose of IL-2 is in the range of from 1-10 MIU daily.
  - 8. The method of claim 5 wherein the cytokine is IFN-a.

- 9. The method of claim 8 wherein the dose of IFN- $\alpha$  is in the range of from 1-10 MIU three times a week.
- 10. The method of any one of claims 1-9 wherein the cytokine is administered in a substantially constant dose during the treatment.
  - 11. The method of any one of claims 1-9 wherein the cytokine is administered in a variable dose during the treatment.
- 10 12. The method of any one of claims 1-11 wherein the cytokine is administered subcutaneously.
  - 13. The method of any one of claims 1-12 wherein the antitumor antibody is selected from antibodies directed against the MN (G250) antigen.
  - 14. The method of any one of claims 1-13 wherein the antitumor antibody is a chimeric or humanized G250 antibody or a fragment thereof.
  - 15. The method of any one of claims 1-14 wherein the antitumor antibody is administered in intervals of from 5-20 days.
- 16. The method of claim 2 wherein the first treatment stage comprises5-20 days.
  - 17. The method of claim 2 wherein the second treatment stage comprises 50-200 days.

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